# Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of the claims in the application.

# Listing of Claims

1. (Currently Amended) A method of causing expression of a desired heterologous protein in gastrointestinal mucosal cells of a mammal, the method comprising placing a nucleotide sequence encoding the desired heterologous protein under the control of a promoter consisting of a nucleotide sequence of SEQ ID NO: 2, the promoter being operatively interconnected to the nucleotide sequence encoding the desired heterologous protein, in a recombinant gut-colonizing bacterium, wherein the recombinant gut-colonizing bacterium is suitably attenuated so that the mammal does not experience significant harmful effects as a result of infection by the recombinant gut-colonizing bacterium, orally administering the recombinant gut-colonizing bacterium to the mammal, and causing expression of the desired heterologous protein in the gastrointestinal mucosal cells of the mammal.

# 2-22. (Cancelled).

 (Previously Presented) The method of claim 1, wherein the desired heterologous protein induces a protective immune response against a pathogen in the mammal.

# 24-25. (Cancelled).

- (Currently Amended) The method of claim 23, 1, wherein the recombinant gut-colonizing gut-colonizing bacterium is a Salmonella spp.
- (Previously Presented) The method of claim 26, wherein the Salmonella spp. is Salmonella typhimurium or Salmonella typhi.

### (Cancelled).

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- (Previously Presented) The method of claim 23, wherein the pathogen is Yersinia pestis.
- (Currently Amended) The method of claim 29, wherein the desired heterologous protein comprises the F1-antigen of Yersinia pestis.
- (Currently Amended) The method of claim 23, wherein the recombinant gut-colonizing gut-colonizing bacterium is administered as a composition which further comprises a pharmaceutically acceptable carrier or diluent.
  - 32. (Cancelled).
- 33. (Currently Amended) A method of inducing a serum or mucosal antibody response in a mammal against Yersinia pestis comprising expressing en F1-antigen of Yersinia pestis in an attenuated recombinant Salmonella spp. by placing a nucleotide sequence encoding the F1-antigen under control of a promoter consisting of e the nucleotide sequence of SEQ ID NO: 2, the promoter being operatively interconnected to the nucleotide sequence, and administering a dosage of the attenuated recombinant Salmonella spp. orally to e the mammal.
- (Previously Presented) The method of Claim 33 wherein the Salmonella spp. is Salmonella typhimurium or Salmonella typhi.
- 35. (Previously Presented) The method of Claim 33 wherein the attenuated recombinant Salmonella spp. is administered with a pharmaceutically acceptable carrier or diluent.